

## **Newborn Screening Report Form**

Infant's Name:	Infant's Birth Date/
Newborn Screening Program Lab #:	_Mother's Name:
[ ] Diagnosis pending, Follow-up Plan:	
Final Diagnosis (please attach CONFIRMATION LAB RE  [] Normal  [] Trait Condition (specify carrier status)  [] Classic Galactosemia (GG phenotype/genotype)  [] Duarte/Galactosemia Compound Heterozygote (DG phenotype)  [] Congenital Hypothyroidism  [] Congenital Adrenal Hyperplasia due to 21-Hydroxylase I  [] Cystic Fibrosis  [] Amino Acid Disorder (specify type)  [] Fatty Acid Oxidation Disorder (specify type)  [] Organic Acid Disorder (specify type)  [] Mucopolysaccharidosis Type I (specify type)  [] Pompe (specify type)  [] Sickle Cell Disease (specify type)  [] Hemoglobin disease (specify type)  [] Biotinidase Deficiency (specify type)  [] Other (specify)	otype/genotype) Deficiency
Treatment Indicated? [ ] yes [ ] no Date treatment started//	
Referred to pediatric sub-specialist (specify name):  [ ] Endocrinologist:  [ ] Metabolic Specialist/Geneticist:  [ ] Neurologist:	Date of first clinic visit to specialist://  [ ] Hematologist:  [ ] Pulmonologist:
Family referred for (check all that apply):  [ ] Genetic counseling (check provider): GeneticistOther (specify)  [ ] Enrollment in Newborn Screening Long-term Follow-u [ ] Early Intervention Services	p Program
Print Physician's Name	Telephone/
Physician Signature	Date//
Mail or Fax this follow-up form with complete diagnostic information and confirmation lab results to: Oklahoma State Department of Health ATTN: Newborn Screening Program Coordinator 123 Robert S. Kerr Ste 1702 Oklahoma City, OK 73102-6406 Fax: (405) 900-7556	Questions or Referral Information Phone: (405) 426-8220 (800) 766-2223